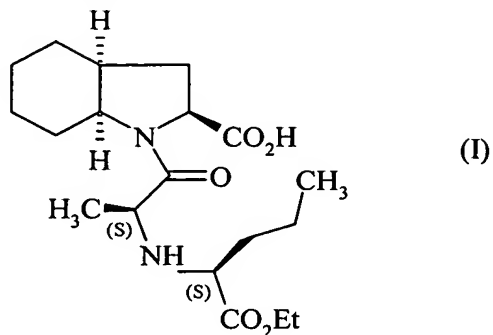


LISTING OF CLAIMS

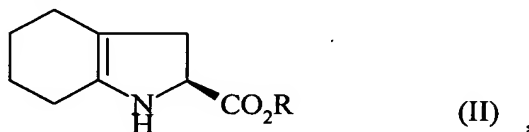
Claims 1-9 (CANCELED)

10. (NEW) A process for the synthesis of a compound of formula (I) :



and its pharmaceutically acceptable salts,

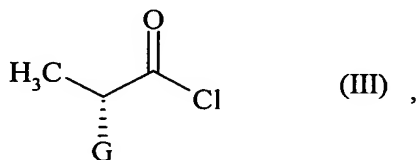
wherein a compound of formula (II) :



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wherein R represents hydrogen, benzyl or linear or branched (C₁-C₆)alkyl, is reacted

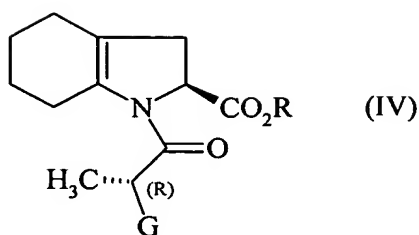
with a compound of formula (III) having the (R) configuration :



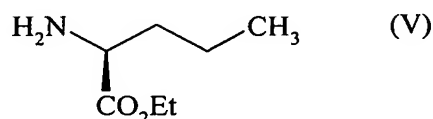
10 wherein G represents chlorine, bromine, iodine, hydroxy, p-toluenesulphonyloxy, methanesulphonyloxy or trifluoromethanesulphonyloxy ,

in the presence of a base,

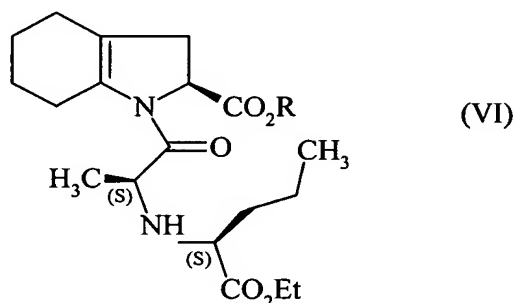
to yield a compound of formula (IV) :



which is reacted with the compound of formula (V) having the (S) configuration :



to yield a compound of formula (VI) :



5

which is hydrogenated in the presence of a catalyst,
to yield, after deprotection where necessary, the compound of formula (I).

- 10 **11. (NEW)** A process according to Claim 10, wherein the base used for the reaction between the compounds of formulae (II) and (III) is an organic amine selected from triethylamine, pyridine and diisopropylethylamine, or a mineral base selected from NaOH, KOH, Na₂CO₃, K₂CO₃, NaHCO₃ and KHCO₃.
- 12. (NEW)** A process according to Claim 10, wherein G represents chlorine, bromine, p-toluenesulphonyloxy, methanesulphonyloxy or trifluoromethanesulphonyloxy.

13. (NEW) A process according to Claim 12, wherein the reaction between the compounds of formulae (IV) and (V) is carried out in the presence of an organic amine selected from triethylamine, pyridine and diisopropylethylamine, or of a mineral base selected from Na_2CO_3 , K_2CO_3 , NaHCO_3 and KHCO_3 .

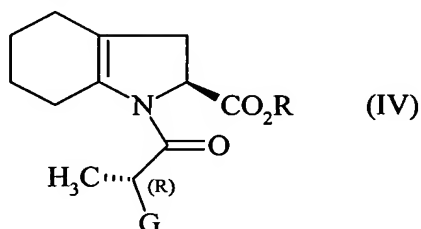
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14. (NEW) A process according to Claim 10, wherein G represents hydroxy .

15. (NEW) A process according to Claim 14, wherein the reaction between the compounds of formulae (IV) and (V) is carried out in the presence of an activation reagent selected from N-methyl-N-phenyl-aminotriphenylphosphonium iodide and hexamethylphosphorus triamide together with ammonium perchlorate, or, when R is other than hydrogen, by Mitsunobu reaction.

16. (NEW) A process according to Claim 10, wherein the catalyst is selected from palladium, platinum, rhodium and nickel.

17. (NEW) A compound selected from those of formula (IV) :



wherein R represents hydrogen, benzyl or linear or branched $(\text{C}_1\text{-C}_6)$ alkyl and G represents chlorine, bromine, hydroxy, p-toluenesulphonyloxy, methanesulphonyloxy or trifluoromethanesulphonyloxy.

10

18. (NEW) A process according to Claim 10 for the synthesis of perindopril in the form of its tert-butylamine salt.